AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

- 1. (Original) A method of effecting cross-linking of a resin comprising generating vinyl sulfonyl moieties *in situ* with the resin, said vinyl sulfonyl moieties then undergoing a reaction which effects cross-linking of the resin.
- 2. (Original) A method as claimed in claim 1 wherein the vinyl sulfonyl moieties are generated as a result of loss of a liquid carrier for the resin to be cross-linked.
- 3. (Original) A method as claimed in claim 2 wherein evaporation of the liquid carrier causes generation of the vinyl sulfonyl moieties.
- 4. (Currently Amended) A method as claimed in any one of claims 1 to 3 wherein cross-linking results from reaction of the vinyl sulfonyl moieties with nucleophilic groups in the resin composition.
 - 5. (Original) A cross-linkable resin composition comprising
 - (i) a polymer to be cross-linked;
 - (ii) a liquid carrier for the polymer;
 - (iii) nucleophilic groups; and
 - (iv) vinyl sulfonyl precursor groups capable of generating vinyl sulfonyl groups on loss of liquid carrier from the composition

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at least one of the groups (iii) and (iv) being attached to the polymer to be cross-linked whereby loss of the liquid carrier results in generation of a vinyl sulfonyl moiety to effect cross-linking of the polymer.

- 6. (Original) A composition as claimed in claim 5 wherein generation of the vinyl sulfonyl moiety results from loss of HX from the vinyl sulfonyl precursor groups, where X is a leaving group.
- 7. (Original) A composition as claimed in claim 6 wherein the vinyl sulfonyl precursor groups are of the formula (II)

where X is a leaving group, and R^1 , R^2 and R^3 are independently selected from a hydrogen atom, a substituted or unsubstituted alkyl group, and a substituted or unsubstituted aryl group, and \sim can be a chemical bond to carbon or heteroatom functionality.

- 8. (Original) A composition as claimed in claim 7 where R¹ is hydrogen.
- 9. (Currently Amended) A composition as claimed in claim 7 or 8-wherein at least one of R² and R³ is hydrogen.
- 10. (Original) A composition as claimed in claim 9 wherein R² and R³ are both hydrogen.
- 11. (Currently Amended) A composition as claimed in any one of claims 6 to 10-wherein the leaving group X is selected from groups of the formula –OR⁴, -OC(O)R⁴,

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 $-NR_2^4$, $-SR^4$, $-NCOOR^4$ or $-OSO_3R^4$ where R^4 is hydrogen, a substituted or unsubstituted alkyl group, or a substituted or unsubstituted aryl group, or X is F, Cl or Br.

- 12. (Original) A composition as claimed in claim 11 wherein X is of the formula $-OR^4$.
 - 13. (Original) A compositions as claimed in claim 12 wherein R⁴ is hydrogen.
- 14. (Original) A composition as claimed in claim 12 wherein R⁴ is methyl or ethyl.
- 15. (Currently Amended) A composition as claimed in any one of claims 6 to 14-wherein the liquid carrier for the polymer to be cross-linked has the formula HX.
- 16. (Currently Amended) A composition as claimed in any one of claims 5 to 15-wherein the vinyl sulfonyl precursor groups are attached to the polymer chains to be cross-linked.
- 17. (Original) A composition as claimed in claim 16 wherein the polymer to be cross-linked comprises 0.5 to 25% by mole of the vinyl sulfonyl precursor groups.
- 18. (Original) A composition as claimed in claim 17 wherein the polymer to be cross-linked comprises 1 to 10% by mole of vinyl sulfonyl precursor groups.
- 19. (Original) A composition as claimed in claim 18 wherein the polymer to be cross-linked comprises 3 to 7% by mole of vinyl sulfonyl precursor groups.

20. (Currently Amended) A composition as claimed in any one of claims 16 to 19-wherein the polymer incorporating the vinyl sulfonyl precursor groups is a co-polymer of a compound of formula (IV)

with other olefinically unsaturated monomers.

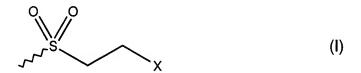
21. (Currently Amended) A composition as claimed in any one of claims 16 to 19 when directly or indirectly dependent from claim 3 wherein the polymer incorporating the vinyl sulfonyl precursor groups is a co-polymer of a compound of formula (V)

in which R^1 - R^3 are as defined in claim 37, R^4 is as defined in claim 711, R^5 , R^6 and R^7 are independently hydrogen or methyl, and n is a positive integer.

22. (Original) A composition as claimed in claim 21 where R⁴ is hydrogen, methyl or ethyl.

- 23. (Currently Amended) A composition as claimed in any one of claims 5 to 22-wherein the nucleophilic groups are selected from –OH, -SH and –NHR⁸ where R⁸ is hydrogen, substituted or unsubstituted alkyl (preferably C₁₋₄ alkyl), or a substituted or unsubstituted aryl, group.
- 24. (Currently Amended) A composition as claimed in any one of claims 5 to 22-wherein the nucleophilic groups are acetoacetoxy groups.
- 25. (Currently Amended) A composition as claimed in any one of claims 5 to 24-wherein the nucleophilic groups are attached to polymer chains to be cross-linked.
- 26. (Original) A composition as claimed in claim 25 wherein the polymer to be cross-linked comprises 0.5 to 25% by mole of the nucleophilic groups.
- 27. (Original) A composition as claimed in claim 26 wherein the polymer to be cross-linked comprises 1 to 10% by mole of the nucleophilic groups.
- 28. (Original) A composition as claimed in claim 19 wherein the polymer to be cross-linked comprises 3 to 7% by mole of the nucleophilic groups.
- 29. (Currently Amended) A composition as claimed in any one of claims 1 to 285 wherein the polymer to be cross-linked is dissolved in the liquid carrier.
- 30. (Currently Amended) A composition as claimed in any one of claims 1-to 285 in the form of a latex comprising a continuous aqueous phase and a discontinuous particulate phase of the polymer to be cross-linked, said polymer having attached thereto said vinyl sulfonyl precursor groups and said nucleophilic groups.

- 31. (Original) A composition as claimed in claim 30 wherein the polymer to be cross-linked has been obtained by copolymerisation of comonomers including vinyl sulfonyl precursor groups, comonomers including nucleophilic groups and optionally additional monomers.
- 32. (Original) A composition as claimed in claim 31 wherein the polymer incorporates said additional monomers which are selected from (meth)acrylic acid, itaconic acid, C₁₋₂₀ (e.g. C₁₋₈) alkyl esters of these acids, vinyl acetate, vinyl versatates, styrene, butadiene, and combinations of the aforesaid monomers.
- 33. (Original) A composition as claimed in claim 32 wherein the additional monomer is selected from vinyl acetate, butyl acrylate, 2-ethylexyl acrylate and butyl methacrylate.
- 34. (Original) A latex comprising a continuous liquid phase and a discontinuous phase of a film-forming polymer incorporating
 - (iii) nucleophilic groups; and
 - (iv) groups of the formula (I)



where X is a leaving groups and $\sim\sim$ represents a chemical bond to a carbon or heteroatom, whereby loss of liquid results in generation of a vinyl sulfonyl moiety by loss of HX on the groups of formula (I) and cross-linking of the polymer by reaction of this vinyl sulfonyl and the nucleophilic groups.

- 35. (Original) An emulsion as claimed in claim 34 wherein the polymer comprises 1 to 10% by mole of the groups of formula (I) and 1 to 10% by mole of the nucleophilic groups.
- 36. (Currently Amended) A latex as claimed in claim 34 or 35 wherein the polymer has been obtained by emulsion polymerisation of
- (A) a compound of formula (IV)

- (B) a hydroxyalkyl (meth)acrylate (e.g. hydroxyethyl acrylate, hydroxyethyl methacrylate, hydroxypropyl acrylate, hydroxypropyl methacrylate), 2-acetoacetoxyethyl acrylate or 2-acetoacetoxyethyl methacrylate; and
- (C) at least one additional monomer.

37. (Original) The compound of formula (IV)

38. (Original) The compound of formula (V)

$$R^{5} \longrightarrow 0 \qquad R^{6} \qquad R^{7} \qquad O \qquad O \qquad R^{2} \qquad R^{3} \qquad (V)$$

in which R^1 - R^3 are as defined in claim 3, R^4 is as defined in claim 7, R^5 , R^6 and R^7 are independently hydrogen or methyl, and n is a positive integer.